

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus.p2n model

Run on: August 23, 2003, 19:55:10 ; Search time 255 Seconds
(without alignments)
3705.111 Million cell updates/sec

Title: US-09-745-506-37
Perfect score: 350
Sequence: 1 MDKALSLINDFASLSFAE.....LENKINILSETDRDFQGVV 350

Scoring table:
OLIGO
Xgapop 60.0 , Xgapext 60.0
Ygapop 60.0 , Ygapext 60.0
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 2552756 seqs, 1349719017 residues

Word size: 1

Total number of hits satisfying chosen parameters: 5103490

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Command line parameters:

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-LOOPEXT=0 -ONITS=bits -START=1 -END=1 -MATRIX=oligo -TRANS=human40.cdi
-LIST=45 -DOCALLIGN=200 -THR_SCORE=quality -THR_MIN=1 -ALIGN=15 -MODE=LOCAL
-OUTFWT=pro -NORR=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US09745506 -ECGN 1.1 0 -runat.22082003.104427.7254 -NCPD=6 -ICPD=3
-NO_MMAP -LARGEORDER -NEG_SCORES=0 -WAIT -DSPBLOCK=100 -LONCLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=60 -XGAPEXT=60 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=60 -YGAPEXT=60 -DELOP=6 -DELEXT=7

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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,

and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	350	100.0	1053	22	AAH52212	Human AFP protein
2	350	100.0	1574	22	AAH16397	Human CDNA sequenc
3	350	100.0	1696	22	AAK60866	Human immune/haema
4	350	100.0	1739	23	ABV23243	Human prostate exp
5	350	100.0	1739	23	ABV29087	Human prostate exp
6	311	88.9	1398	22	AAFS9945	Human gene express
7	296	84.6	1554	22	AA544644	Human full-length
8	211	60.3	1385	24	ABF60919	Human protein kina
9	155	44.3	1686	23	AA585172	DNA encoding novel
10	122	34.9	462	22	AAU23953	Human breast cance
11	122	34.9	796	22	AAH07192	Human CDNA clone (
12	119	35.0	14969	22	AAK78763	Human immune/haema
13	90	25.7	514	22	AAI15105	Human breast cell
14	73	20.9	465	22	ABA46423	Human breast cell
15	73	20.9	465	22	ABA57019	Human foetal liver
16	73	20.9	465	22	AAK05073	Human brain expres
17	73	20.9	465	22	AAI15235	Probe #5168 for ge
18	73	20.9	465	22	AAI04973	Probe #4964 used t
19	73	20.9	465	23	ABS30297	Human liver single
20	70	20.0	249	21	AA25260	Human secreted pro
21	69	19.7	208	22	ABA51524	Human breast cell
22	69	19.7	208	22	ABA69581	Human foetal liver
23	69	19.7	208	22	AAK17792	Human brain expres
24	69	19.7	208	22	AAI24414	Probe #14347 for g
25	69	19.7	208	22	AAI09950	Probe #9941 used t
26	69	19.7	208	23	ABS43283	Human liver single
27	53	15.1	394	25	ABA45683	Bovine EST associa
28	49	14.0	633	22	AA534287	Human CDNA encodin
29	43	12.3	453	23	AA544816	Human contig polyn
30	38	10.9	515	22	AA585169	DNA encoding novel
31	25	7.1	522	23	AA585171	DNA encoding novel
32	19	5.4	60	24	ABN33267	Human spliced tren
33	19	5.4	273	23	ABV35380	Human prostate exp
34	19	5.4	273	23	ABV44212	Human prostate exp
35	19	5.4	443	23	ABV14293	Human prostate exp
36	18	5.1	357	23	ABV05124	Human prostate exp
37	13	3.7	41	24	ABL60924	Human protein kina
38	10	2.9	339	21	AAI17249	Human secreted pro
39	10	2.9	400	18	AAV75090	Staphylococcus aur
40	10	2.9	463	22	AAK63571	Human immune/haema
41	10	2.9	1083	22	AAH53378	S. epidermidis ope
42	10	2.9	1104	24	ABN91378	Staphylococcus epi
43	10	2.9	3041	22	AAH55004	S. epidermidis gen
44	10	2.9	3138	22	AAH54589	S. epidermidis gen
45	10	2.9	3441	22	AAH54443	S. epidermidis gen

ALIGNMENTS

RESULT 1
AAH52212
ID AAH52212 standard; CDNA; 1053 BP.
AC AAH52212;
XX
DT 10-SEP-2001 (first entry)
XX
DE Human AFP protein encoding CDNA sequence SEQ ID NO:239.
XX
KW Human; secreted protein; secretion; bacterial cell; fungal cell;
KW eukaryotic cell; fusion protein; maltose binding protein;
KW immunoglobulin constant region; polynucleotide tag; ss.
XX
OS Homo sapiens.
XX
PN WO200129221-A2.

XX Primer sets for synthesizing polynucleotides, particularly the 5602
 PT full-length cDNAs defined in the specification, and for the detection
 PT and/or diagnosis of the abnormality of the proteins encoded by the
 PT full-length cDNAs -

PS Claim 8: SEQ ID 15359: 2537pp + CD ROM; English.

XX The present invention describes primer sets for synthesizing 5602
 CC full-length cDNAs defined in the specification. Where a primer set
 CC comprises: (a) an oligo-of primer and an oligonucleotide complementary
 CC to the complementary strand of a polynucleotide which comprises one of
 CC the 5602 nucleotide sequences defined in the specification, where the
 CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
 CC of an oligonucleotide comprising a sequence complementary to the
 CC complementary strand of a polynucleotide which comprises a 5'-end
 CC sequence and an oligonucleotide comprising a sequence complementary to a
 CC polynucleotide which comprises a 3'-end sequence, where the
 CC oligonucleotide comprises at least 15 nucleotides and the combination of
 CC the 5'-end sequence/3'-end sequence is selected from those defined in
 CC the specification. The primer sets can be used in antisense therapy and
 CC in gene therapy. The primers are useful for synthesizing polynucleotides,
 CC particularly full-length cDNAs. The primers are also useful for the
 CC detection and/or diagnosis of the abnormality of the proteins encoded by
 CC the full-length cDNAs. The primers allow obtaining of the full-length
 CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
 CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
 CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
 CC represent oligonucleotides, all of which are used in the exemplification
 CC of the present invention.

XX SQ Sequence 1574 BP; 420 A; 361 C; 372 G; 421 T; 0 other;

Alignment Scores:

Pred. No.: 0 Length: 1574
 Score: 350.00 Matches: 350
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 22 Gaps: 0

US-09-745-506-37 (1-350) x AAH16397 (1-1574)

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 DB 271 ATGGATTGTAAGGCTCTCTTCTCTTGAATGATTTGCAATCCCTCTGCTTGAG 330
 OY 21 SerTPAspAsnValGlyLeuLeuValGluProSerProHisThrValAsnThrLeu 40
 DB 331 AGTTGGACAAATGGTGGATTACTGCGGAACCAACCCACATCACTGTAATACACTC 390
 OY 41 PheLeuThrAsnAspLeuThrGluGluValMetGluGluValLeuGlnLysLysAlaAsp 60
 DB 391 TTCCTGACCAATGACTGCTGACTGAGGAAGTGAAGAGAGTGTGAAAAAGAGGAGAC 450
 OY 61 LeuLeuLeuSerTyrisProPheArgProMetLysArgLleThrTPAsnThr 80
 DB 451 CTCATCTCTCTCTACATCCGCTATCTTCGACCCATGAAGGCACTTAACCTGGAACACA 510
 OY 81 TPPLysGluArgLeuValIleArgAlaLeuGluAsnArgValGlyLysSerProHis 100
 DB 511 TGGAAAGACCGCTGGTATCCGGGCTCGAAGACAGAGTGTGCTCTCTCCAT 570
 OY 101 ThrAlaTyraAspAlaAlaProGlnGlyValAsnAsnTrpLeuAlaLysGlyLeuGlyAla 120
 DB 571 ACAGCTATGATGCTGCGCCCAAGGGCGTCAACACTGTTGGCTAAAGGCTTGAGCT 630
 OY 121 CysThrSerArgProIleHisProSerLysAlaProAsnTyxProThrGluGlyAsnHis 140
 DB 631 TGTACTCCAGGCGCATCATCTTCCAAAGCTCCCACTACCTTCACAGAGGAAACACAC 650
 OY 141 ArgValGluPheAsnValAsnTyxThrGlnAspLeuAspLysValMetSerAlaValLys 160

DB 691 CGAGTAGAATTCACGTTACTACACCAAGACCTGGACAAAGTCACTGCTGACGTGAAA 750
 OY 161 GlyTlaAspGlyValSerValThrSerPheSerAlaArgThrGlyLysGlnGluThr 180
 DB 751 GGAAATGAGAGGTGTTCTCTGCTCACTCTTTTCTGCTAGGACTGTAATGAGAAACAA 810
 OY 181 ArgIlaAsnLeuAsnGlyThrGlnLysAlaLeuMetGlnValValAspPheLeuSerArg 200
 DB 811 CGGATTAATCTGATTTGTTCTGCTGAGAGCTTTGATGAGAGTGTGATTTCTTCCGG 870
 OY 201 AsnLysGlnLeuTyrisGlnLysThrGluIleLeuSerLeuGluLysProLeuLeuHis 220
 DB 871 AACAAACAATTATCAGAAAGACGAAATCTGTCACTGGAGAAAGCTTGTCTTCAAT 930
 OY 221 ThrGlyMetGlyArgLeuGlySerThrLeuAspGluSerValSerLeuAlaThrMetIleAsp 240
 DB 931 ACAGGAATGAGAGGTTATGACACATGGAATGATCTGTCTCCCTGGCAACCATGATTTGAT 990
 OY 241 ArgIlaLysArgHisLeuLysLeuSerHisIleArgLeuAlaLeuGlyValGlyArgThr 260
 DB 991 CGAATTAATAAGACACCTTAATAACTATCTCATATTCGCTTACCCCTTGGGGTGGAGAAC 1050
 OY 261 LeuGluSerGlnValLysValAlaAlaLeuGlyAlaGlySerGlySerValLeuGln 280
 DB 1051 TTGAGCTCTCAAGTCAAAAGTCGTGGCCGTGTGCTGTCTGAGAGACGCTTGACAG 1110
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 OY 321 SerAspLeuArgAspMetLeuAspSerHisLeuGluAsnLysIleAsnIleIleLeuSer 340
 DB 1231 TCTGACCTTCGAAATATGCTGAGATCTCACTTGGAGAAATGAATATATATATCTATCA 1290
 OY 341 GluThrAspArgAspProLeuGlnValVal 350
 DB 1291 GAGACTGACAGGAGCCCTCTTCAGGTGGTA 1320

RESULT 3
 AAK60866
 ID AAK60866 standard; cDNA; 1696 BP.
 AC AAK60866;
 XX 06-NOV-2001 (first entry)
 DE Human Immune/haematopoietic antigen encoding cDNA SPO ID NO:5926.
 XX KW Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
 XX cytosolic; gene therapy; vaccine; metastasis; ss.
 OS Homo sapiens.
 XX MO200157182-A2.
 PD 09-AUG-2001.
 PF 17-JAN-2001; 2001MO-US01354.
 XX 31-JAN-2000; 2000US-0179065.
 PR 04-FEB-2000; 2000US-0180628.
 PR 24-FEB-2000; 2000US-0184664.
 PR 02-MAR-2000; 2000US-0186350.
 PR 16-MAR-2000; 2000US-0189874.
 PR 17-MAR-2000; 2000US-0190076.
 PR 18-APR-2000; 2000US-0198123.
 PR 19-MAY-2000; 2000US-0205513.
 PR 07-JUN-2000; 2000US-0209467.
 PR 28-JUN-2000; 2000US-0214886.

PR 30-JUN-2000; 2000US-0215135.
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PR 05-DEC-2000; 2000US-0251030.
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PR 08-DEC-2000; 2000US-0251856.
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PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-0259678.

(HUMA-) HUMAN GENOME SCI INC.
PA Rosen CA, Barash SC, Ruben SM;
PI WPI; 2001-483426/52.
XX P-PSDB; AAM88085.
XX
XX Nucleic acids encoding human Immune/hematopoietic antigen polypeptides,
PT useful for preventing, diagnosing and/or treating cancers and
PT metastasis -
XX
XX
XX Claim 1; SEQ ID NO 5926; 3071pp + Sequence Listing; English.
XX
XX AAK54951 to AAK64702 encode the human Immune/haematopoietic antigen (I)
XX amino acid sequences given in AAM82170 to AAM91921. (I) have cytostratic
XX activity, and can be used in gene therapy and vaccine production. (I)
XX proteins and polynucleotides may be used in the prevention, diagnosis and
XX treatment of diseases associated with inappropriate (I) expression. For
XX example, they may be used to treat disorders associated with decreased


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Db      296 AGTGGGACAAATTGGATTACTGTGGGAACCAAGCCACCATCTACTGTAATACACTC 355
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Db      356 TTCTGTGACCAATGACCGATGAGGAAGATGATGAGAGAGGTGCTGCAAAAGAGGACGAC 415
Oy      61  LeuIleLeuSerLysHisProProlIlePheArgProMetLysArgIle-ThrTrpAsnTh 80
Db      416 CTCATTCTCTCCATCCATCCGCTATCTTCCAGCCCTGAAAGGCGCATTAACCTGGAAAC 475
Oy      80  rTTP-Lys-GluArgLeuValIleArgAlaLeuGlnAsnArgValGlyIleTyrSerPro 99
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Oy      100 HisThrAlaTyrAspAlaAlaProGlnGlyValAsnAsnTrpLeuAlaLysGlyLeuGly 119
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Db      596 GCTGTGACCTCCAGGCCCATCATCTTCCAAAGCTCCAACTACCTACAGAGGAGAAC 655
Oy      140 HisArgValGluPheAsnValAsnTyrThrGlnAspLeuAspLysValMetSerAlaVal 159
Db      656 CACCGAGTAGAATTCACAGCTTAACTACACCAAGACCTGGACAAGTCATGTCTGCAAGTG 715
Oy      160 LysGlyIleAspGlyValSerValThSerPheSerAlaArgThrGlyAsnGluGln 179
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Db      776 ACACGGATTATCTGAATGTACTCAGAGGCTTGTGATGCGAGTGATGATTTCTTTC 835
Oy      200 ArgAsnLysGlnLeuTyrGlnLysThrGluIleLeuSerLeuGlnLysProLeuLeu 219
Db      836 CGGAACAACAACCTTATCAGAGACGGAATTCGTGTCACGAGAAACCTTGGCTCTA 895
Oy      220 HisThrGlyMetGlyArgLeuCysThrLeuAspGluSerValSerLeuAlaThrMetIle 239
Db      896 CATACTGGAAATGGGCGGTGTATGCACACTGGATGATGTCTCTCCCTGGCAACCATGATT 955
Oy      240 AspArgIleLysArgHisLeuLysLeuSerHisIleArgLeuAlaLeuGlyValGlyArg 259
Db      956 GATCGAATTAACAAAGACCTTAACATCTCATATCCCTTAAGCCCTTGGGGTGGGAGA 1015
Oy      260 ThrLeuGlnSerGlnValLysValValAlaLeuCysAlaGlySerGlySerValLeu 279
Db      1016 ACCTTAGAGTCTCAAGTCAAGAGTCGTGCCCTGTGCTGCTGGTGGAGACGCTTCTG 1075
Oy      280 GlnGlyValGluAlaAspLeuTyrLeuThrGlyGluMetSerHisHisAspThrLeuAsp 299
Db      1076 CAGGGTGTGGAGGCTGACCTTATCTCAGAGGTGAGATGCTCCATCATGATATCTTGGAT 1135
Oy      300 AlaAlaSerGlnGlyIleAsnValIleLeuCysGlnHisSerAsnThrGluArgLysPhe 319
Db      1136 GCTGCTTCCCAAGAAATTAATGCTCTCTGTGAAACAGCAACACTGACGAGAGCTTT 1195
Oy      320 LeuSerAspLeuArgAspMetLeuAspSerHisLeuGlnLysLysIleAsnIleLeuLeu 339
Db      1196 CTTTCTGACCTTCGAGATATGCTGATCTCTCACTTGGAGAAATATATATATTCCTA 1255
Oy      340 SerGluThrAspArgAspProLeuGlnValVal 350
Db      1256 TCAGAGACTGACAGGAGACCTCTTCAGGGGTGA 1288

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RESULT 7
AAS44644
ID      AAS44644 standard; DNA; 1554 BP.
XX      AAS44644;
AC      AAS44644;
XX      18-DEC-2001 (first entry)
DT

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```

XX      Human full-length polynucleotide sequence #69.
DE
XX      Mammal; human; rhesus monkey; baker's yeast; fission yeast; Norway rat;
KW      mouse; Chinese hamster; African clawed frog; fruit fly; dog; leukaemia;
KW      cancer; lymphoma; neuroblastoma; autoimmune disorder; cell proliferation;
KW      nervous system disorder; inflammatory disorder; cell differentiation; ds;
KW      angiogenesis; stem cell growth factor; activin; inhibin; cartilage; burn;
KW      genetic disorder; bone regeneration; tendon; ligament; tissue repair;
KW      antihypertensive; immunosuppressive; vasotropic; antiparkinsonian;
KW      neuroprotective; osteopathic; antidiabetic; antistimulant; antiallergic;
KW      immunostimulant; analgesic; gene therapy.
XX
OS      Homo sapiens.
XX
PN      WO200164834-A2.
XX
PD      07-SEP-2001.
XX
PF      26-FEB-2001; 2001WO-US04926.
XX
PR      28-FEB-2000; 2000US-0515126.
PR      18-MAY-2000; 2000US-0577409.
PR      17-JUN-2000; 2000US-0597707.
PR      14-JUL-2000; 2000US-0616807.
PR      19-SEP-2000; 2000US-0664641.
XX
PA      (HXSE-) HXSEQ INC.
XX
PI      Tang YT, Liu C, Zhou P, Asundi V, Zhang J, Zhao QA, Ren F;
PI      Xue AJ, Yang Y, Wehrman T, Wang J, Ma Y, Wang D, Chen R, Xu C;
PI      Dmanac R;
XX
DR      WPI: 2001-589862/66.
XX
DR      P-PSDB; AAU27744.
XX
PT      Novel polypeptides and nucleic acids obtained from cDNA libraries
PT      prepared from various human tissues, for diagnosis, treatment of
PT      cancer, neurological, inflammatory disorders and for use in arrays for
PT      detection.
XX
PS      Claim 1; SEQ ID NO 69; 153bp; English.
XX
CC      Sequences AAS44576-AAS44919 represent full-length polynucleotides and
CC      config polynucleotides encoding polypeptides of the invention. The DNA
CC      and protein sequences are useful for the treatment, diagnosis and
CC      prevention of various types of disorder in a mammalian subject such as a
CC      human, dog, monkey, mouse, hamster or rat. The disorders include cancers
CC      such as leukemia, lymphoma and neuroblastoma, autoimmune disorders such
CC      as multiple sclerosis, connective tissue disease, rheumatoid arthritis,
CC      diabetes mellitus, allergic rhinitis, asthma and eczema, nervous system
CC      disorders such as Parkinson's disease, Alzheimer's disease, Huntington's
CC      chorea, amyotrophic lateral sclerosis, spinal muscular atrophy and
CC      Wernicke disease, inflammatory disorders such as nephritis, Crohn's
CC      disease, ischemia-reperfusion injury, shock, sepsis and inflammatory
CC      bowel disease. The sequences exhibit activity relating to angiogenesis,
CC      cell proliferation, cell differentiation, stem cell growth factor,
CC      activin or inhibin. Therefore, they can be used to manipulate stem cells
CC      in culture to give rise to neuroepithelial cells that can be used to
CC      augment or replace cells damaged by illness, accidental damage or genetic
CC      disorders. The sequences may also be used for regeneration of bone,
CC      cartilage, tendons and ligaments and in tissue repair and burn healing.
CC      Note: Some sequences for this patent did not form part of the printed
CC      specification, but were obtained in electronic format directly from WIPO
CC      at ftp.wipo.int/pub/published_pcl_sequences.
XX
SQ      Sequence 1554 BP; 428 A; 358 C; 346 G; 422 T; 0 other;

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Alignment Scores:
Pred. No.:      2,67e-290      Length:      1554
Score:          296.00          Matches:      349
Percent Similarity: 99.43%      Conservative: 0

```


Best Local Similarity: 99.438
 Query Match: 84.578
 DB: 22 Gaps: 0
 US-09-745-506-37 (1-350) x AAS44644 (1-1554)

OY 1 MetAspLeuLysAlaLeuLeuSerSerLeuAsnAspPheAlaSerLeuSerPheAlaGlu 20
 DB 236 ATGGAATTGAAGGCTCTCTTCTTCTTCTGAAATGACTTGGACCTCCCTCCTTGGCTGAG 295
 OY 21 SerTTPAspAsnValGlyLeuLeuValGluProSerProProHisThrValAsnThrLeu 40
 DB 296 AGTTGGACAAATGGTGGATTACTGGTGGAACCAAGCCACACATCTGTAATATACATC 355
 OY 41 PheLeuThrAsnAspLeuThrGluGluValMetGluGluValLeuGlnLysLysAlaAsp 60
 DB 356 TTCTCCAGCAATGACTGCTGAGAGAAAGATGAGAGAGCTGCTGAAAAAGAGGACAGAC 415
 OY 61 LeuLeuLeuSerThrHisProProIlePheArgProMetLysArgIleThrTPAsnThr 80
 DB 416 CTCATTCTCTCTACCATCCGCTATCTTCCGACCCATGAGCGCATTAACCTGGAACACA 475
 OY 81 TPPLysGluArgLeuValIleArgAlaLeuGluAsnArgValGlyIleThrSerProHis 100
 DB 476 TGGAAAGAGCGCGCTGGTATCCGGGCTCGAGAAACAGAGTGGTATCTACTCTCCTCAT 535
 OY 101 ThrAlaTPAspAlaAlaProGlnGlyValAsnAsnTPLeuAlaLysGlyLeuGlyAla 120
 DB 536 ACAGCCTATGATGCTGCGCCGACGAGGCGTCAACACTGGTGGTAAAGCGCTTGGAGCT 595
 OY 121 CysThrSerArgProIleHisProSerLysAlaProAsnTPYrProThrGluGlyAsnHis 140
 DB 596 TGTACTCTCCAGGCCATCATCTTCCCAAGCTCCCACTACCTCCACAGAGGAACACAC 655
 OY 141 ArgValGluPheAsnValAsnTPYrThrGlnAspLeuAspLysValMetSerAlaValLys 160
 DB 656 CGAGTAAGAATTCACCTTACTACACCCAAAGACTGGACAAAGTCTGCTGCAAGTGA 715
 OY 161 GlyIleAspGlyValSerValThrSerPheSerAlaArgThrGlyAsnGluGluGlnThr 180
 DB 716 GGAATTGACGGTGTCTCTCTCTCTCTTCTTCTGCTGAGACTGGTATGAGAACAAACA 775
 OY 181 ArgIleAsnLeuAsnCysThrGlnLysAlaLeuMetGlnValValAspPheLeuSerArg 200
 DB 776 CGGAATTAATCTGAATGTACTCAGAAAGCTTTGATGACAGTGGTAAATTTCTTCCCG 835
 OY 201 AsnLysGlnLeuTPYrGlnLysThrGluIleLeuSerLeuGluLysProLeuLeuHis 220
 DB 836 AACCAACAACTTATCTCAGAAAGCGGAAATCTGTCTGCTGAGAACGCTTGGCTTACAT 895
 OY 221 ThrGlyMetGlyArgLeuCysThrLeuAspGluSerValSerLeuAlaThrMetIleAsp 240
 DB 896 ACTGGAATGGACGGTATATGACACACTGGATGAATCTGTCTCCCTGGCAACATGATGAT 955
 OY 241 ArgIleLysArgHisLeuLysLeuSerHisIleArgLeuAlaLeuGlyValGlyArgThr 260
 DB 956 CGAATAAAAAGACACTAAACTATCTCATATTCGTTAGCCCTTGGGGTGGGGAACACC 1015
 OY 261 LeuGluSerGlnValLysValValAlaLeuCysAlaGlySerGlySerSerValLeuGln 280
 DB 1016 TTAGAGTCTCAAGTCAAAAGTCTGGCCCTGTGTGCTGCTGGGAGCAACGCTTCGAC 1075
 OY 281 GlyValGluAlaAspLeuTPYrLeuThrGluValMetSerHisHisAspThrLeuAspAl 300
 DB 1076 GGTGTGGAGCGTACCTTAACTCACAGAGTGAATGTCCCATCATATATAT-TTTGGATGC 1134
 OY 300 AlaSerGlnGlyIleAsnValIleLeuCysGluHisSerAsnThrGluArgGlyPheLe 320
 DB 1135 TCGTCCCAAGAAATATGTCATCTCTGTGAACACACACACACAGAGGAGGCTTCT 1194
 OY 320 userAspLeuArgAspMetLeuAspSerHisLeuGluAsnLysIleAsnIleLeuSe 340
 DB 1195 TTCTGACCTTCGAGATATGCTGGAATTCCTGAGATTAAGATAAATATATCTCTATC 1254

OY 340 rgiuThrAspArgAspProLeuGlnValVal 350
 DB 1255 AGAGACTGACAGGAGCCCTCTTCAAGTGGTA 1285

RESULT 8

ABL60919
 ID ABL60919 standard; cDNA; 1385 BP.
 AC ABL60919;

23-SEP-2002 (first entry)

Human protein kinase C 27.17 polypeptide encoding cDNA.

Human; protein kinase C 27.17; protein metabolism; gene; ss.

OS Homo sapiens.

FH Key Location/Qualifiers
 FT CDS 389..1132

/*tag= a /product= "protein kinase C 27.17 polypeptide"

CN1333355-A.

30-JUN-2002.

07-JUL-2000; 2000CN-0117049.

07-JUL-2000; 2000CN-0117049.

(SHAN-) SHANGHAI BIODOOR GENE DEV CO LTD.

Mao Y, Xie Y;

WPI; 2002-305609/35.

P-PSDB; ABB08182.

Human protein kinase C 27.17 polypeptide and its encoding polynucleotide, for treating e.g. protein metabolism disturbance -

Claim 6; Page 25-26 (disclosure); 33pp; Chinese.

CC The invention relates to a human protein kinase C 27.17 polypeptide and
 CC its encoding polynucleotide. The polypeptide can be expressed by standard
 CC DNA recombination. The polynucleotide, polypeptide and its antagonist are
 CC useful for treating e.g. protein metabolism disturbance. The present
 CC sequence represents the human protein kinase C 27.17 polypeptide encoding
 CC cDNA.

SQ Sequence 1385 BP; 375 A; 324 C; 308 G; 378 T; 0 other;

Alignment Scores:

Pred. No.: 4.5e-204 Length: 1385
 Score: 211.00 Matches: 346
 Percent Similarity: 98.02% Conservative: 0
 Best Local Similarity: 98.02% Mismatches: 4
 Query Match: 60.29% Indels: 7
 DB: 24 Gaps: 0

US-09-745-506-37 (1-350) x ABL60919 (1-1385)

OY 1 MetAspLeuLysAlaLeuLeuSerSerLeuAsnAspPheAlaSerLeuSerPheAlaGlu 20
 DB 82 ATGGAATTGAAGGCTCTCTTCTTCTTCTGAAATGACTTGGACCTCCCTCCTTGGCTGAG 141
 OY 21 SerTTPAspAsnValGlyLeuLeuValGluProSerProProHisThrValAsnThrLeu 40
 DB 142 AGTTGGACAAATGGTGGATTACTGGTGGAACCAAGCCACACATCTGTAATATACATC 201
 OY 41 PheLeuThrAsnAspLeuThrGluGluValMetGluGluValLeuGlnLysLysAlaAsp 60

Db 202 TTCCTGACCAATGACCTGACTGAGGAAGTGTGAGGAGAGGTGCTGCAAAAGACGCCAC 261
Qy 61 LeuileuSerTyRHISProProIlePheArProMetLysArgIleThrPAsnThr 80
Db 262 CTATTTCCTGCTACATCCGCGCTATCTTCGACCACTGAAAGCCATTAACCTGGAACACA 321
Qy 81 TrpLysGluArgLeuValIleArgAlaLeuGluAsnArgValGlyIleTyrSerProHis 100
Db 322 TGGAAAGAGCGCCGCGGTGATCCGGCTGTGAGAAAGAGCGGTATCTACTCTCCAT 381
Qy 101 ThrAlaTyrAspAlaAlaProGlnGlyValAsnAsnTrpLeuAlaLysGlyLeuGlyAla 120
Db 382 ACAGCCTATGATGTGGGCCGCCAGGCGCTCAACAACGTGGCTTAAGGCGCTGGAGCT 441
Qy 121 CysThrSerArgProIleHisProSerLysAlaProAsnTyr--ProThrGluLysAsn- 139
Db 442 TGTACCTCCAGGCC-ATACATCTTCCAA-CCGCCAACCTT-CCCCACAGAGG--AACCC 496
Qy 140 HisArgValGluPheAsnValAsnTyrThrGlnAspLeuAspLysValMetSerAlaVal 159
Db 497 CACGAGTGAATTCACAGCTTAACCTACACCAAGACCTGGACAAAGTCATGTCTGCACTG 556
Qy 160 LysGlyIleAspGlyValSerValThrSerPheSerAlaArgThrGlyAsnGluGluGln 179
Db 557 AAAGGAATTGACGGTGTCTCTGCACTTCTTTCTGCTAGAGCTGGTAAATGAGGACAA 616
Qy 180 ThrArgIleAsnLeuAsnCysThrGlnLysAlaLeuMetGlnValAlaAspPheLeuSer 199
Db 617 ACAGGGAATTATTCGAATTGTACACAGAGCTTTGATGCAAGGTGTGATTTCTTCC 676
Qy 200 ArgAsnLysGlnLeuTyrGlnLysThrGluIleLeuSerLeuGluLysProLeuLeu 219
Db 677 CGGACAAACAACCTTATCAGAAACAGGAAATTCGTGTCAGGAAACCTTTGCTTCTA 736
Qy 220 HisThrGlyMetGlyArgLeuCysThrLeuAspGluSerValSerLeuAlaThrMetIle 239
Db 737 CATATCGAATGGAGCGGTATTCACACATCGATGATGTCTCCCTGGCAACATGATTT 796
Qy 240 AspArgIleLysArgHisLeuLysLeuSerHisIleArgLeuAlaLeuGlyValGlyArg 259
Db 797 GATCGAATTAAGAACACCTTAACCTATTCATATTCCTTACAGCCCTTGGGGGAGAGA 856
Qy 260 ThrLeuGluSerGlnValLysValValAlaLeuCysAlaGlySerGlySerSerValLeu 279
Db 857 ACCTTAGAGTCTCAAGTCAAGTCGCGGCTGTGTGTGTGTGTGGAGACGCTTGTG 916
Qy 280 GlnGlyValGluAlaAspLeuTyrLeuThrGlyLysMetSerHisHisAspThrLeuAsp 299
Db 917 CAGGGTGTGGAGCGTACCTTACTCTCACAGGTGAGATGTCCTCATCATGATCTTTGGAT 976
Qy 300 AlaAlaSerGlnGlyIleAsnValIleLeuCysGluHisSerAsnThrGluArgGlyPhe 319
Db 977 GCTGCTCCCAAGAAATAATGTCATCTGTGAAACAGACACAGACAGAGAGCTTT 1036
Qy 320 LeuSerAspLeuArgAspMetLeuAspSerHisIleGluAsnLysIleAsnIleLeu 339
Db 1037 CTTTCGACCTTCGAGATATGTGATTCCTCACTTGAGAAATTAATTTATCTCA 1096
Qy 340 SerGluThrAspArgAspProLeuGlnValVal 350
Db 1097 TCAGAGACTGACAGGAGCCCTTTCAGGTGTA 1129

RESULT 9
ID AAS85172/c
AAS85172 standard; cDNA; 1686 BP.
XX AAS85172;
DT 13-FEB-2002 (first entry)
DE DNA encoding novel human diagnostic protein #20976.
XX Human; chromosome mapping; gene mapping; gene therapy; forensic;

KM food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX Homo sapiens.
OS WO200175067-A2.
FN 11-OCT-2001.
PD 30-MAR-2001; 2001WO-US08631.
PF 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX (HYSE-) HYSEQ INC.
PA Drmanac RT, Liu C, Tang YT;
PI WPI: 2001-639362/73.
DR P-PSDB: ABG20985.
XX New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
XX Claim 1: SEQ ID No 20976; 103pp; English.
PS The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
SQ Sequence 1686 BP; 445 A; 397 C; 391 G; 452 T; 1 other;

Alignment Scores:
Pred. No.: 3.78e-147 Length: 1686
Score: 155.00 Matches: 194
Percent Similarity: 98.48% Conservative: 0
Best Local Similarity: 98.48% Mismatches: 0
Query Match: 44.29% Indels: 3
DB: Gaps: 0

US-09-745-506-37 (1-350) x AAS85172 (1-1686)

Qy 82 LysGluArgLeuValIleArgAlaLeuGluAsnArgValGlyIleTyrSerProHisThr 101
Db 1115 AAGGAGCGCGCTGTATCCGCGCTGTGAGAAACAGTCGTGATCTACTCTCTCATTTACA 1056
Qy 102 AlaTyrAspAlaAlaProGlnGlyValAsnAsnTrpLeuAlaLysGlyLeuGlyAlaCys 121
Db 1055 GCCTATGATGCTGCCGCCAGGCGCTCAACAACGTGGCTTAAGGCGCTTGGAGCTTGT 996
Qy 122 ThrSerArgProIleHisProSerLysAlaProAsnTyrProThrGluLysHisArg 141
Db 995 ACCTCCAGGCCATCATCTCTTCCAAAGCTCCCACTACCTTACAGAGGAAACCAACCGCA 936

QY	142	ValGILPheAsnValAsnThrYthrGlnAspLeu-AspLysValMetSer-ALAValLysG	161
Db	935	GTFGAATTCACTTACTACTACCCCAAGACCTGGAGCAAGTCATGCTGGAGTGAAG	87/67
QY	161	LYtle-AspGlyValSerValThrSerPheSerAlaArgThrGlyAsnGluGlnThr	180
Db	875	GAATTTCAGAGGGTTCCTCTCACTCTTTCTGCTAGACTGGTAAGAGAAACA	81/6
QY	181	ArgIleAsnLeuAsnCysThrGlnLysAlaLeuMetGlnValValAspPheLeuSerArg	200
Db	815	CGGATTAACTGCAATGTTACTCAGAAAGCCTTTCAGACGAGGTGATATTCTTCCCGG	75/6
QY	201	AspLysGlnLeuThrGlnLysThrGluIleLeuSerLeuGluLysProLeuLeuHis	220
Db	755	AACAAACAACCTTATTCAGAAAGACGAAATCTGTCTACAGGAAAGCCCTTCTTCAT	69/6
QY	221	ThrGlyMetGlyArgLeuCysThrLeuAspGlnSerValSerLeuAlaThrMetIleAsp	240
Db	695	ACTGGAATGGAGCGGTATGACACACTGGATGAAATGTCTCCCTGGCAACATATGAT	63/6
QY	241	ArgIleLysArgHisIleuLysLeuSerHisIleArgLeuAlaLeuGlyValGlyArgThr	260
Db	635	CGAATTAAGAACACCTAAACATATCTATATTCGCTTAGCCCTGGGGTGGAGAAC	57/67
QY	261	LeuGlnSerGlnValLysValValAlaLeuCysAlaGlySerGly	275
Db	575	TTAGACTTCAGACTCAAAATCGTGGCCCTGTGTGCTGTCTGGG	531
RESULT 10			
AL23953	ID	AA23953 standard; cDNA; 462 BP.	
XX	XX	AA23953:	
XX	AC	07-DEC-2001 (first entry)	
XX	DT	Human breast cancer expressed polynucleotide 16410.	
XX	XX	Human: breast cancer; cell marker; cytostatic; ss.	
XX	OS	Homo sapiens.	
XX	PN	WO200151628-A2.	
XX	PD	19-JUL-2001.	
XX	PF	10-JAN-2001; 2001WO-US00798.	
XX	PR	14-JAN-2000; 2000US-0176077.	
XX	PR	14-MAR-2000; 2000US-0189167.	
XX	PR	24-MAR-2000; 2000US-0192099.	
XX	PR	29-MAR-2000; 2000US-0193480.	
XX	PR	15-MAY-2000; 2000US-0205230.	
XX	PR	09-JUN-2000; 2000US-0211315.	
XX	PR	25-JUL-2000; 2000US-0220534.	
XX	PA	(MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.	
XX	PI	Lillie J, Xu Y, Wang Y, Steinmann K;	
XX	DR	WPI; 2001-451856/48.	
XX	PS	New peptide useful as a marker for the diagnosis of breast cancer	
XX	PS	Claim 1; Page 3004; 3695pp; English.	
XX	CC	The invention relates to human breast cancer expressed polynucleotides	
XX	CC	(AA207544-AA26789) and methods of assessing whether a patient is	
XX	CC	afflicted with breast cancer by examining the correlation between the	
XX	CC	expression of certain markers and the cancerous state of breast cells.	
XX	CC	The polynucleotides and encoded polypeptides are potential markers for	
XX	CC	detecting, diagnosing, monitoring, characterizing, treating and	
XX	CC	potentially preventing breast cancer. The polynucleotides and encoded	

CC	polypeptides are also useful for isolating compounds with cytostatic activity.			
CC				
XX				
SQ	Sequence 462 BP; 103 A; 126 C; 118 G; 115 T; 0 other;			
Alignment Scores:				
	Pred. No.:	3,34e-114	Length:	462
	Score:	122.00	Matches:	122
	Percent Similarity:	100.00%	Conservative:	0
	Best Local Similarity:	100.00%	Mismatches:	0
	Query Match:	34.86%	Indels:	0
DB:		22	Gaps:	0
US-09-745-506-37 (1-350) x AAL23953 (1-462)				
QY	1 MetAspleuYsAlaLeuLeuSerSerLeuAsnAspPheAlaSerLeuSerPheAlaGlu 20			
Db	96 ATGGATTGTAAGGCTCTCTTCTTCTTCTTGATGACTTTGGATCCCTCTGTTGGCGAG 155			
QY	21 SerTrpAspAsnValIcylLeuLeuValGluProSerProProHisThrValAsnThrLeu 40			
Db	156 AGTTGGGACAAATTGATTACTGTGGGAAACCAACCCACACACTACTAAATACATC 215			
QY	41 PheLeuThrAsnAspleuthrGluGluValMetGluGluValIleGlnIcylsAlaAsp 60			
Db	216 TTCTTGACCAATATACCTTGACTGAGAGAGATGATGAGAGGTGCTGCANNAAGCCAGAC 275			
QY	61 LeuIleLeuSerTrhAspProIlePheArgPrometLysArgIleThrTrpAsnThr 80			
Db	276 CTCATTCTCTCCACCATCGCGCTATCTTCGACCCAGACCAAGAGCGATTAACCTGGACACA 335			
QY	81 TrpLysGluArgLeuValIleArgAlaLeuGluAsnArgValcylIleTrpSerProHis 100			
Db	336 TGGAGAGAGCGCTCGTGTGATCCGGGCTGTGGAGAACAGAGTCGGTATCTACTCTTCAT 395			
QY	101 ThrAlaTrpAspAlaIleProGlnGluValAsnAsnTrpLeuAlaIcylLeuGluValA 120			
Db	396 ACAGCCATATATCTCTCGGCCCGAGGCGTCACCAACTGTTGGCTAAAGSGCTTGAGCT 455			
QY	121 CysThr 122			
Db	456 TGTACC 461			
RESULT 11				
ID	AAH07192 standard; cDNA; 796 BP.			
XX	AAH07192;			
XX	AAH07192;			
XX	26-JUN-2001 (first entry)			
DT				
XX	Human cDNA clone (5'-primer) SEQ ID NO:4027.			
DE				
XX	Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss			
XX	Homo sapiens.			
OS	EPI074617-A2.			
PN	07-FEB-2001.			
XX				
PD				
XX	28-JUL-2000; 2000EP-0116126.			
PF				
XX	29-JUL-1999; 99JP-0248036.			
PR	27-AUG-1999; 99JP-0300253.			
PR	11-JAN-2000; 2000JP-0118776.			
PR	02-MAY-2000; 2000JP-0183767.			
PR	09-JUN-2000; 2000JP-0241899.			
XX				
PA	(HELI-) HELIX RES INSTR.			
XX				
XX	Ota T, Tsogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;			
PI	Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;			

XX WPI; 2001-318749/34.
 XX Primer sets for synthesizing polynucleotides, particularly the 5602
 PT full-length cDNAs defined in the specification, and for the detection
 PT and/or diagnosis of the abnormality of the proteins encoded by the
 PT full-length cDNAs -
 XX
 PS Claim 1; SEQ ID 4027; 2537bp + CD ROM; English.

XX The present invention describes primer sets for synthesizing 5602
 CC full-length cDNAs defined in the specification. Where a primer set
 CC comprises: (a) an oligo-dt primer and an oligonucleotide complementary
 CC to the complementary strand of a polynucleotide which comprises one of
 CC the 5602 nucleotide sequences defined in the specification, where the
 CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
 CC of an oligonucleotide comprising a sequence complementary to the
 CC complementary strand of a polynucleotide which comprises a 5'-end
 CC sequence and an oligonucleotide comprising a sequence complementary to a
 CC polynucleotide which comprises a 3'-end sequence, where the
 CC oligonucleotide comprises at least 15 nucleotides and the combination of
 CC the 5'-end sequence/3'-end sequence is selected from those defined in
 CC the specification. The primer sets can be used in antisense therapy and
 CC in gene therapy. The primers are useful for synthesizing polynucleotides,
 CC particularly full-length cDNAs. The primers are also useful for the
 CC detection and/or diagnosis of the abnormality of the proteins encoded by
 CC the full-length cDNAs. The primers allow obtaining of the full-length
 CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
 CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
 CC AAB95893 represent human amino acid sequences; and AAH13632
 CC represent oligonucleotides, all of which are used in the exemplification
 CC of the present invention.
 CC
 XX SQ Sequence 796 BP; 192 A; 204 C; 202 G; 195 T; 3 other;

Alignment Scores:
 Pred. NO.: 5.68e-114 Length: 796
 Score: 122.00 Matches: 122
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 34.86% Indels: 0
 DB: Gaps: 0

US-09-745-506-37 (1-350) x AAH07192 (1-796)

QY 1 MetaspLeuYsAlaLeuLeuSerSerLeuAsnAspPheAlaSerLeuSerPheAlaGlu 20
 DB 271 ATGATTTGAGAGGCTCTCTCTTCTTGAATGACCTTGCATCCCTCGTTGCTGAG 330
 QY 21 SerTrpAspAsnValGlyLeuLeuValGluProSerProProHisThrValAsnThrLeu 40
 DB 331 AGTTGGGACATGTTGGATTGCTGGTGAACCAACCCACACACATGTAATACACTC 390
 QY 41 PheLeuThrAsnAspLeuThrGluGluValMetGluGluValLeuGluLysLysAlaAsp 60
 DB 391 TTCTGACCATGACCTGACTGAGAGACTGATGGAGGGGTGCTCAAAAGAGCAAC 450
 QY 61 LeuLeuLeuSerTyrHisProProIlePheArgProMetLysArgIleThrTrpAsnThr 80
 DB 451 CTCATTTCTCTCTCCATCCGCTGCTATCTTCGACCATGAGCGCATTAACCTGAACACA 510
 QY 81 TrpLysGluArgLeuValIleArgAlaLeuGluAsnArgValGlyIleTyrSerProHis 100
 DB 511 TGGAGAGGCGCTGGTATCCGGGCTCTGGAGAACAGTCGATATCTACTCTCTCAT 570
 QY 101 ThrAlaTyrAspAlaAlaProGlnGlyValAsnAsnTrpLeuAlaLysGlyLeuGlyAla 120
 DB 571 AAGGCTTATGATGCTGCGCCCAAGGCTCAACAACATGTTGGCTAAAGGGCTTGAGCT 630
 QY 121 CysThr 122
 DB 631 TGTACC 636

RESULT 12
 ID AAK78763
 AC AAK78763; standard; DNA; 14969 BP.
 XX
 DT 07-NOV-2001 (first entry)
 XX
 DE Human Immune/haematopoietic antigen genomic sequence SEQ ID NO:33575.
 DE
 KW Human; Immune; haematopoietic; immune/haematopoietic antigen; cancer;
 KW Cystostatic; gene therapy; vaccine; metastasis; ds.
 XX
 OS Homo sapiens.
 XX
 PN WO200157182-A2.
 PD
 XX 09-AUG-2001.
 XX
 FE 17-JAN-2001; 2001WO-US01354.
 XX
 PR 31-JAN-2000; 2000US-0179065.
 PR 04-FEB-2000; 2000US-0180628.
 PR 24-FEB-2000; 2000US-0184664.
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 PR 08-DEC-2000; 2000US-0251989.
 PR 08-DEC-2000; 2000US-0251990.
 PR 11-DEC-2000; 2000US-0254097.
 PR 05-JAN-2001; 2001US-0259678.
 XX
 PA (HUMA-) HUMAN GENOME SCI INC.
 PI Rosen CA, Barash SC, Ruben SM;
 XX
 DR WPI; 2001-483426/52.
 XX
 PT Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
 PT useful for preventing, diagnosing and/or treating cancers and
 PT metastasis
 XX
 PS Disclosure; SEQ ID NO 33575; 3071pp + Sequence Listing; English.
 XX
 CC AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)
 CC amino acid sequences given in AAK82170 to AAK91921. (I) have cytostatic
 CC activity, and can be used in gene therapy and vaccine production. (I)
 CC proteins and polynucleotides may be used in the prevention, diagnosis and
 CC treatment of diseases associated with inappropriate (I) expression. For
 CC example, they may be used to treat disorders associated with decreased
 CC expression by rectifying mutations or deletions in a patient's genome
 CC that affect the activity of (I) by expressing inactive proteins or to
 CC supplement the patients own production of (I). Additionally, (I)
 CC polynucleotides may be used to produce the secreted (I), by inserting
 CC the nucleic acids into a host cell and culturing the cell to express the
 CC protein. (I) proteins and polynucleotides may be used to prevent,
 CC diagnose and treat immune/hematopoietic-related diseases, especially
 CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703
 CC to AAK87694 represent human immune/hematopoietic antigen genomic
 CC sequences from the present invention. AAK54942 to AAK54950 and AAK82169
 CC represent sequences used in the exemplification of the present invention.
 XX
 SQ Sequence 14969 BP; 4016 A; 3069 C; 3042 G; 4842 T; 0 other;
 Alignment Scores:
 Pred. No.: 1 11e-109 Length: 14969
 Score: 119.00 Matches: 119
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 34.00% Indels: 0
 DB: 22 Gaps: 0
 US-09-745-506-37 (1-350) x AAK78763 (1-14969)
 QY 1 MetAspLeuYsAlaLeuSerSerLeuAsnAspPheAlaSerLeuSerPheAlaGlu 20
 DB 2604 ATGGATTGGAAGGCTCCTCTTCTCTTGAATACCTTTGCATCCCTCGTTGCTGAG 2603
 QY 21 SerTrpAspAsnValGlyLeuLeuValGluProSerProProHisThrValAsnThrLeu 40
 DB 2664 AGTTGGACATGTTGGATTACTGTGGAACCAAGCCACACATGTAATAATACCTC 2723
 QY 41 PheLeuThrAsnAspLeuThrGluValMetCysGluValLeuGlnIbysLyAlaAsp 60
 DB 2724 TTCCTGACCAATGACCTGAGTGAAGAGTGTGAGAGGCTGCAAAAGAGCGAGAC 2783
 QY 61 LeuIleLeuSerTrpHisProProIlePheArgProMetLeuYsArgIleThrTrpAsnThr 80
 DB 2784 CTCATTCTCTCTACCACTCCGCCCTATCTTCGACCCATGAAGCCATRAACTGGAACA 2843
 QY 81 TrpLySGluArgLeuValIleArgAlaLeuGluAsnArgValGlyIleYrSerProHis 100
 DB 2844 TGGAAAGAGCGCGTGGATCCGGCTGTGAGAAACAGAGTACGATCTACTCTCTCAT 2903

QY 101 ThrAlaTrAspAlaAlaProGlnGlyValAsnAsnTrpLeuAlaGlyGly 119
|||||
Db 2904 ACAGCCATGATGCTGCGCCCGAGGCGTCACACACTGGTTGGCTAAAGGCTTGGT 2960

RESULT 13
AL15105
ID AL15105 standard; cDNA; 514 BP.
XX
AC AL15105;
XX
DF 07-DEC-2001 (first entry)
XX
DE Human breast cancer expressed polynucleotide 7562.
XX
KW Human; breast cancer; cell marker; cytostatic; ss.
XX
OS Homo sapiens.
XX
PN WO200151628-A2.
XX
PD 19-JUL-2001.
XX
PF 10-JAN-2001; 2001WO-US00798.
XX
PR 14-JAN-2000; 2000US-0176077.
PR 14-MAR-2000; 2000US-0189167.
PR 24-MAR-2000; 2000US-0192099.
PR 29-MAR-2000; 2000US-0193480.
PR 15-MAY-2000; 2000US-0205230.
PR 09-JUN-2000; 2000US-0211315.
PR 25-JUL-2000; 2000US-0220534.
XX
PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
PI Little J, Xu Y, Wang Y, Steinmann K;
PI WPI: 2001-451856/48.
XX
DR New peptide useful as a marker for the diagnosis of breast cancer -
XX
PT Claim 1; Page 1361; 3695pp; English.
XX
PS The invention relates to human breast cancer expressed polynucleotides
XX CC (AA107544-AA126789) and methods of assessing whether a patient is
CC afflicted with breast cancer by examining the correlation between the
CC expression of certain markers and the cancerous state of breast cells.
CC The polynucleotides and encoded polypeptides are potential markers for
CC detecting, diagnosing, monitoring, characterizing treating and
CC potentially preventing breast cancer. The polynucleotides and encoded
CC polypeptides are also useful for isolating compounds with cytostatic
CC activity.
XX
SQ Sequence 514 BP; 113 A; 141 C; 130 G; 126 T; 4 other;

Alignment Scores:
Pred. No.: 1.12e-81 Length: 514
Score: 90.00 Matches: 90
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 25.71% Indels: 0
Gaps: 0

US-09-745-506-37 (1-350) x AL15105 (1-514)

QY 1 MetAspLeuYsAlaLeuLeuSerLeuAsnAspPheAlaSerLeuSerPheAlaGlu 20
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Db 122 ATGGATTGAAGCGCTCTCTTCCTGATGACTTGGCATCCCTCTCGTTGGCTAG 181

QY 21 SerTrpAspAsnValGlyLeuLeuValGluProSerProTrpOHisTrpValAsnThrLeu 40
|||||
Db 182 AGTTGGACAATGTTGATTAATCTGTTGGAACCAAGCCCGACATACACTTAATACACTC 241

QY 41 PheLeuThrAsnAspLeuThrGluGluValMetGluGluValLeuGlnTyrAlaAsp 60
|||||
Db 242 TTCTTGACCAATGACTGCTGACTGAGAGTGATGAGAGAGTGCCTGCAAAAAGGACGAC 301

QY 61 LeuLeuLeuSerTrpHisProProIlePheArgProMetLysArgIleTrpPasnThr 80
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Db 302 CTCATTCCTCCCTACATCCGCGCTATCTTCGAGCCCAATGAAGCGCATACCTGGACACA 361

QY 81 TrpLysGluArgLeuValIleArgAlaLeu 90
|||||
Db 362 TGGAGAGAGCGCCTGTGATCCGGGCTCTG 391

RESULT 14
ABA46423/C
ID ABA46423 standard; DNA; 465 BP.
XX
AC ABA46423;
XX
DF 01-FEB-2002 (first entry)
XX
DE Human breast cell single exon nucleic acid probe #5118.
XX
KW Human; microarray; single exon probe; gene expression; breast;
KW disease; cancer; ss.
XX
OS Homo sapiens.
XX
PN WO200157271-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00662.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
PI Penn SG, Hanzel DK, Chen W, Rank DR;
PI WPI: 2001-496933/54.
XX
DR New spatially-addressable set of single exon nucleic acid probes,
XX useful for measuring gene expression in sample derived from human
XX breast, comprises number of single exon nucleic acid probes -
XX
PT Claim 1; SEQ ID NO 5118; 327pp + sequence listing; English.
XX
PS The invention relates to a spatially-addressable set of single exon
XX CC nucleic acid probes for measuring gene expression in a sample derived
XX CC from human breast and B7 474 cells. The method involves contacting
XX CC the probes with a collection of detectably labelled nucleic acids
XX CC derived from mRNA of human breast, and then measuring the label
XX CC bound to each probe of the microarray. The probes are useful for
XX CC verifying the expression of regions of genomic DNA predicted to
XX CC encode proteins. They are useful for gene discovery, and for
XX CC determining predisposition and/or prognosing breast disease. Gene
XX CC expression analysis is useful for assessing the toxicity of chemical
XX CC agents on cells. The microarray of this invention presents a far greater
XX CC diversity of probes for measuring gene expression, with far less bias
XX CC than expressed sequence tag microarrays. The method is suitable for
XX CC rapid production of functional information from genomic sequence. The
XX CC present sequence is a single exon nucleic acid probe of the invention.
XX CC Note: The sequence data for this patent did not form part of the
XX CC printed specification, but was obtained in electronic format directly
XX CC from WIPO at http://wipo.int/pubd/published_pct_sequences.

Sequence 465 BP; 138 A; 109 C; 93 G; 125 T; 0 other;

Alignment Scores:

Pred. No.:	1,83e-64	Length:	465
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Query Match:	20.86%	Indels:	0
DB:	22	Gaps:	0

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OY 47 ThrGIUGluValMetGIUGluValLeuGlnLysLysAlaAspLeuIleLeuSerTyrHis 66
DB 463 ACTGAGGAAGTATGAGGAGGCTGCTGCAAAAGAGGACAGCTCATCTCTCTACCAT 404
OY 67 ProProlIlePheArgProMetLysArgIleThrTyrAsnThrTyrPylsGIUArgLeuVal 86
DB 403 CCGCTATCTTCCGACCCATGAGCGCATTAACCTGGACACATGGAAGAGACGCCCTGGTG 344
OY 87 IleArgAlaLeuGluAsnArgValGlyIleTyrSerProHisThrAlaTyrAspAlaAla 106
DB 343 ATCCGGGCTCTGGAGACAGACTCGGTATCTACTCTCCATACAGCCTATATGATGCTGG 284
OY 107 ProGInGIyValAsnAsnTrpLeuAlaLysGIyLeuGIy 119
DB 283 CCCAGGGCGTCACAACTGTTGGCTAAAGGGCTTGGT 245
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RESULT 15

ABA57019/c

ID ABA57019 standard; DNA; 465 BP.

XX ABA57019;

XX 01-FEB-2002 (first entry)

XX Human foetal liver single exon nucleic acid probe #5324.

XX Human; foetal liver; gene expression; single exon nucleic acid probe; ss.

XX Homo sapiens.

XX WO200157277-A2.

XX 09-AUG-2001.

XX 30-JAN-2001; 2001WO-US00669.

XX 04-FEB-2000; 2000US-0180312.

XX 26-MAY-2000; 2000US-0207456.

XX 30-JUN-2000; 2000US-0608408.

XX 03-AUG-2000; 2000US-0632366.

XX 21-SEP-2000; 2000US-0234687.

XX 27-SEP-2000; 2000US-0236359.

XX 04-OCT-2000; 2000GB-0024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

XX Penn SG, Hanzel DK, Chen W, Rank DR;

XX WPI; 2001-483447/52.

XX Human genome-derived single exon nucleic acid probes useful for

XX analyzing gene expression in human fetal liver -

XX Claim 1; SEQ ID NO 5324; 639bp + sequence listing; English.

XX The invention relates to a single exon nucleic acid probe for

XX measuring human gene expression in a sample derived from human foetal

XX liver. The single exon nucleic acid probes may be used for predicting,

XX measuring and displaying gene expression in samples derived from human

XX fetal liver. The present sequence is a single exon nucleic acid

XX probe of the invention.

XX Note: The sequence data for this patent did not form part of the

CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 465 BP; 138 A; 109 C; 93 G; 125 T; 0 other;

Alignment Scores:

Pred. No.:	1,83e-64	Length:	465
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Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	20.86%	Indels:	0
DB:	22	Gaps:	0

US-09-745-506-37 (1-350) x ABA57019 (1-465)

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DB 463 ACTGAGGAAGTATGAGGAGGCTGCTGCAAAAGAGGACAGCTCATCTCTCTACCAT 404
OY 67 ProProlIlePheArgProMetLysArgIleThrTyrAsnThrTyrPylsGIUArgLeuVal 86
DB 403 CCGCTATCTTCCGACCCATGAGCGCATTAACCTGGACACATGGAAGAGACGCCCTGGTG 344
OY 87 IleArgAlaLeuGluAsnArgValGlyIleTyrSerProHisThrAlaTyrAspAlaAla 106
DB 343 ATCCGGGCTCTGGAGACAGACTCGGTATCTACTCTCCATACAGCCTATATGATGCTGG 284
OY 107 ProGInGIyValAsnAsnTrpLeuAlaLysGIyLeuGIy 119
DB 283 CCCAGGGCGTCACAACTGTTGGCTAAAGGGCTTGGT 245
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